

SKYCLARYS- omaveloxolone capsule

Reata Pharmaceuticals, Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use SKYCLARYS safely and effectively. See full prescribing information for SKYCLARYS.

SKYCLARYS™ (omaveloxolone) capsules, for oral use

Initial U.S. Approval: 2023

INDICATIONS AND USAGE

SKYCLARYS is indicated for the treatment of Friedreich's ataxia in adults and adolescents aged 16 years and older. (1)

DOSAGE AND ADMINISTRATION

- Obtain alanine aminotransferase (ALT), aspartate aminotransferase (AST), bilirubin, B-type natriuretic peptide (BNP), and lipid parameters prior to initiating SKYCLARYS and during treatment. (2.1, 5.1, 5.2, 5.3)
- Recommended dosage is 150 mg (3 capsules) taken orally once daily. (2.2)
- Administer SKYCLARYS on an empty stomach at least 1 hour before eating. (2.2)
- Swallow SKYCLARYS capsules whole. Do not open, crush or chew. (2.2)
- Moderate and Severe Hepatic Impairment: The recommended dosage of SKYCLARYS is 100 mg once daily for patients with moderate hepatic impairment. If adverse reactions emerge, further reduce the dosage to 50 mg once daily. Avoid use in patients with severe hepatic impairment. (2.5, 8.6, 12.3)

DOSAGE FORMS AND STRENGTHS

Capsules: 50 mg (3)

CONTRAINDICATIONS

None. (4)

WARNINGS AND PRECAUTIONS

- Elevation of Aminotransferases: Monitor ALT, AST, and total bilirubin prior to initiation, every month for the first 3 months of treatment, and periodically thereafter. (2.1, 5.1)
- Elevation of B-type Natriuretic Peptide (BNP): Advise patients of signs and symptoms of fluid overload. (2.1, 5.2)
- Lipid Abnormalities: Monitor cholesterol periodically during treatment. (2.1, 5.3)

ADVERSE REACTIONS

Most common adverse reactions (incidence $\geq 20\%$ and greater than placebo) are elevated liver enzymes (AST/ALT), headache, nausea, abdominal pain, fatigue, diarrhea, and musculoskeletal pain. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Reata Pharmaceuticals, Inc. at 1-800-314-3934 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Moderate or Strong CYP3A4 Inhibitors: Avoid concomitant use. Consider SKYCLARYS dosage reduction with monitoring if use is unavoidable. (2.4, 7.1)
- Moderate or Strong CYP3A4 Inducers: Avoid concomitant use. (7.1)

-----**USE IN SPECIFIC POPULATIONS**-----

Pregnancy: Based on animal data, may cause fetal harm. (8.1)

See 17 for **PATIENT COUNSELING INFORMATION** and FDA-approved patient labeling.

Revised: 2/2023

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

SKYCLARYS is indicated for the treatment of Friedreich's ataxia in adults and adolescents aged 16 years and older.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Testing Before Initiating SKYCLARYS and Monitoring to Assess Safety

Obtain ALT, AST, bilirubin, BNP, and lipid parameters prior to initiating SKYCLARYS and during treatment [see *Warnings and Precautions* (5.1, 5.2, 5.3)].

2.2 Recommended Dosage

The recommended dosage of SKYCLARYS is 150 mg (3 capsules) taken orally once daily.

- Administer SKYCLARYS on an empty stomach at least one hour before eating [see *Clinical Pharmacology* (12.3)].
- Swallow SKYCLARYS capsules whole. Do not open, crush, or chew.

2.3 Missed Doses

If a dose of SKYCLARYS is missed, take the next dose at its scheduled time the following day. A double dose should not be taken to make up for a missed dose.

2.4 Recommendations for Concomitant Use with Strong or Moderate CYP3A4 Inhibitors and Inducers

The recommended dosage for concomitant use of SKYCLARYS with cytochrome P450 (CYP) 3A4 inhibitors and inducers are described in Table 1 [see *Drug Interactions* (7.1) and *Clinical Pharmacology* (12.3)].

Table 1: Recommended Dosage of SKYCLARYS with Concomitant Use of CYP3A4 Inhibitors and Inducers

Concomitant Drug Class	Dosage
Strong CYP3A4 inhibitor	<p>Recommended to avoid concomitant use.</p> <p>If coadministration cannot be avoided:</p> <ul style="list-style-type: none">• Reduce the dosage of SKYCLARYS to 50 mg once daily with close monitoring for adverse reactions.

	<ul style="list-style-type: none"> If adverse reactions emerge, coadministration with strong CYP3A4 inhibitors should be discontinued.
Moderate CYP3A4 inhibitor	<p>Recommended to avoid concomitant use.</p> <p>If coadministration cannot be avoided:</p> <ul style="list-style-type: none"> Reduce the dosage of SKYCLARYS to 100 mg once daily with close monitoring for adverse reactions. If adverse reactions emerge, further reduce the dosage of SKYCLARYS to 50 mg once daily.
Strong or Moderate CYP3A4 inducer	Recommended to avoid concomitant use.

2.5 Recommended Dosage for Patients with Hepatic Impairment

The recommended dosage for patients with hepatic impairment are described in Table 2 [see *Use in Specific Populations (8.6)*].

Table 2: Recommended Dosage in Patients with Hepatic Impairment

Impairment Classification (Child-Pugh)	Dosage
Severe (Child-Pugh Class C)	Avoid use
Moderate (Child-Pugh Class B)	<ul style="list-style-type: none"> 100 mg once daily with close monitoring for adverse reactions Consider lowering to 50 mg once daily if adverse reactions emerge
Mild (Child-Pugh Class A)	150 mg once daily

3 DOSAGE FORMS AND STRENGTHS

SKYCLARYS capsules contain 50 mg of omaveloxolone, and are supplied as opaque hard capsules having a light green body and blue cap, imprinted with “RTA 408” in white ink on the body and “50” in white ink on the cap.

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Elevation of Aminotransferases

Treatment with SKYCLARYS can cause an elevation in hepatic transaminases (ALT and AST). In Study 1 [see *Clinical Studies (14)*], the incidence of elevations of ALT or

AST above 5 times and 3 times the upper limit of normal (ULN) was 16% and 31%, respectively, in patients treated with SKYCLARYS. There were no cases of concomitant elevation of transaminases and total bilirubin observed in Study 1. Maximum increases in ALT and AST occurred within 12 weeks after starting SKYCLARYS. Increases in serum aminotransferases were generally asymptomatic and reversible following discontinuation of SKYCLARYS. Patients with clinically significant liver disease were excluded from Study 1.

Monitor ALT, AST, and total bilirubin prior to initiation of SKYCLARYS, every month for the first 3 months of treatment, and periodically thereafter. If transaminases increase to levels greater than 5 times the ULN, or greater than 3 times the ULN with evidence of liver dysfunction (e.g., elevated bilirubin), immediately discontinue SKYCLARYS and repeat liver function tests as soon as possible. If transaminase levels stabilize or resolve, SKYCLARYS may be reinitiated with an appropriate increased frequency of monitoring of liver function [*see Adverse Reactions (6.1) and Use in Specific Populations (8.6)*].

5.2 Elevation of B-Type Natriuretic Peptide

Treatment with SKYCLARYS can cause an increase in BNP, a marker of cardiac function. In Study 1, a total of 14% of patients treated with SKYCLARYS had an increase from baseline in BNP and a BNP above the ULN (100 pg/mL), compared to 4% of patients who received placebo. The incidence of elevation of BNP above 200 pg/mL was 4% in patients treated with SKYCLARYS. Cardiomyopathy and cardiac failure are common in patients with Friedreich's ataxia. Patients were excluded from Study 1 if they had BNP levels > 200 pg/mL prior to study entry, or a history of clinically significant left-sided heart disease and/or clinically significant cardiac disease, with the exception of mild to moderate cardiomyopathy associated with Friedreich's ataxia [*see Adverse Reactions (6.1)*]. Whether the elevations in BNP in Study 1 are related to SKYCLARYS or cardiac disease associated with Friedreich's ataxia is unclear.

Elevations in BNP may indicate cardiac failure and should prompt an evaluation of cardiac function. Check BNP prior to initiation of SKYCLARYS. Monitor patients for the signs and symptoms of fluid overload, such as sudden weight gain (3 pounds or more of weight gain in one day, or 5 pounds or more of weight gain in a week), peripheral edema, palpitations, and shortness of breath. If signs and symptoms of fluid overload develop, worsen, or require hospitalization, evaluate BNP and cardiac function, and manage appropriately. Management of fluid overload and heart failure may require discontinuation of SKYCLARYS.

5.3 Lipid Abnormalities

Treatment with SKYCLARYS can cause changes in cholesterol. In Study 1, 29% of patients treated with SKYCLARYS reported elevated cholesterol above ULN at one or more time points. Mean increases were observed within 2 weeks of initiation of SKYCLARYS and returned to baseline within 4 weeks of discontinuing treatment. A total of 16% of patients treated with SKYCLARYS had an increase in low-density lipoprotein cholesterol (LDL-C) from baseline, compared to 8% of patients who received placebo.

The mean increase in LDL-C for all SKYCLARYS-treated patients was 23.5 mg/dL at 48 weeks. A total of 6% of patients treated with SKYCLARYS had decreases in high-density lipoprotein cholesterol (HDL-C) from baseline compared to 4% of patients who received placebo. The mean decrease in HDL-C for all SKYCLARYS-treated patients was 5.3 mg/dL at 48 weeks.

Assess lipid parameters prior to initiation of SKYCLARYS and monitor periodically during treatment. Manage lipid abnormalities according to clinical guidelines.

6 ADVERSE REACTIONS

The following clinically significant adverse reactions are described in greater detail in other labeling sections:

- Elevation of aminotransferases [see *Warnings and Precautions (5.1)*]
- Elevation of BNP [see *Warnings and Precautions (5.2)*]
- Lipid abnormalities [see *Warnings and Precautions (5.3)*]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared with rates in clinical trials of another drug and may not reflect the rates observed in clinical practice.

The safety of SKYCLARYS 150 mg once daily has been evaluated in 165 patients with Friedreich's ataxia, including 137 patients exposed for at least 48 weeks, and 125 patients exposed for at least 96 weeks.

The most common adverse reactions in Study 1 ($\geq 20\%$ and greater than placebo) were elevated liver enzymes (AST/ALT), headache, nausea, abdominal pain, fatigue, diarrhea, and musculoskeletal pain. Table 3 shows the adverse reactions that occurred in 10% or more of patients treated with SKYCLARYS and greater than placebo.

Table 3 Adverse Reactions Reported in 10% or More of Patients Treated with SKYCLARYS and Greater than Placebo (Study 1)

Adverse Reactions	SKYCLARYS 150 mg (N = 51) %	Placebo (N = 52) %
Elevated liver enzymes (AST/ALT)	37	2
Headache	37	25
Nausea	33	13
Abdominal pain	29	6
Fatigue	24	14
Diarrhea	20	10
Musculoskeletal pain	20	15
Oropharyngeal pain	18	6
Influenza	16	6
Vomiting	16	12
Muscle spasms	14	6
Back pain	13	8
Decreased appetite	12	4
Rash	10	4

Laboratory Abnormalities

In addition to elevated liver enzymes, additional laboratory abnormalities include elevation of BNP and lipid abnormalities [see *Warnings and Precautions (5.1, 5.2, 5.3)*].

7 DRUG INTERACTIONS

7.1 Effect of Other Drugs on SKYCLARYS

CYP3A4 Inhibitors

Omaveloxolone is a CYP3A4 substrate. Concomitant use of SKYCLARYS with moderate or strong CYP3A4 inhibitors is expected to result in clinically significant increased exposure of omaveloxolone [see *Clinical Pharmacology (12.3)*], which may increase the risk of adverse reactions. Avoid concomitant use of SKYCLARYS with moderate or strong CYP3A4 inhibitors. If use cannot be avoided, dosage modifications are recommended [see *Dosage and Administration (2.4)*].

CYP3A4 Inducers

Omaveloxolone is a CYP3A4 substrate. Concomitant use of SKYCLARYS with moderate or strong CYP3A4 inducers may significantly decrease exposure of omaveloxolone [see *Clinical Pharmacology (12.3)*], which may reduce the effectiveness of SKYCLARYS. Avoid concomitant use of SKYCLARYS with moderate or strong CYP3A4 inducers.

7.2 Effect of SKYCLARYS on Other Drugs

CYP3A4 and CYP2C8 Substrates

Omaveloxolone is a weak inducer of CYP3A4 and CYP2C8. Concomitant use with SKYCLARYS can reduce the exposure of CYP3A4 and CYP2C8 substrates which may reduce the activity of these substrates [see *Clinical Pharmacology (12.3)*]. Refer to the prescribing information of substrates of CYP3A4 and CYP2C8 for dosing instructions if used concomitantly with SKYCLARYS and monitor for lack of efficacy of the concomitant treatment.

Hormonal Contraceptives

Omaveloxolone is a weak CYP3A4 inducer [see *Clinical Pharmacology (12.3)*]. Concomitant use with SKYCLARYS may reduce the efficacy of hormonal contraceptives. Advise patients to avoid concomitant use with combined hormonal contraceptives (e.g., pill, patch, ring), implants, and progestin only pills [see *Use in Specific Populations (8.3)*].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate data on the developmental risks associated with the use of SKYCLARYS in pregnant women. In animal studies, administration of omaveloxolone during pregnancy or throughout pregnancy and lactation produced evidence of

developmental toxicity (embryofetal mortality and growth impairment, and mortality, growth impairment, and neurobehavioral deficits in offspring) at plasma exposures similar to or less than exposures in humans (see Animal Data). In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively. The background risk of major birth defects and miscarriage for the indicated population is unknown.

Data

Animal Data

Oral administration of omaveloxolone (0, 1, 3, or 10 mg/kg/day) to pregnant rats throughout organogenesis resulted in no adverse effects on embryofetal development; however, in a dose range-finding study, oral administration of omaveloxolone at doses up to 30 mg/kg/day to pregnant rats throughout organogenesis produced increases in post-implantation loss and resorptions, resulting in a decrease in viable fetuses, and reduced fetal weight at the highest dose tested. At the highest dose tested in the pivotal study (10 mg/kg/day), plasma exposure (AUC) was approximately 5 times that in humans at the recommended human dose (RHD) of 150 mg/day.

Oral administration of omaveloxolone (0, 3, 10, or 30 mg/kg/day) to pregnant rabbits throughout organogenesis resulted in increased embryofetal mortality and skeletal variations and reduced fetal weight at the highest dose tested, which was associated with maternal toxicity. At the no-effect dose for adverse effects on embryofetal development (10 mg/kg/day), plasma exposure was less than that in humans at the RHD.

Oral administration of omaveloxolone (0, 1, 3, or 10 mg/kg/day) to rats throughout pregnancy and lactation resulted in an increase in stillbirths and impaired neurobehavioral function (increased locomotor activity and learning and memory deficits) in offspring at all doses, reduced body weight in offspring at all but the lowest dose tested, and delayed sexual maturation (males), increased postnatal mortality, and impaired reproductive performance in offspring at the highest dose tested. A no-effect dose for adverse effects on pre- and postnatal development was not identified. Plasma exposure (AUC) at the lowest dose tested was less than that in humans at the RHD.

8.2 Lactation

Risk Summary

There are no data on the presence of omaveloxolone or its metabolites in human milk. The effects on milk production and the breastfed infant are unknown. Omaveloxolone was excreted in the milk of lactating rats following oral administration. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for SKYCLARYS and any potential adverse effects on the breastfed infant from SKYCLARYS or from the underlying maternal condition.

8.3 Females and Males of Reproductive Potential

SKYCLARYS may decrease the efficacy of hormonal contraceptives [see *Drug Interactions (7.2) and Clinical Pharmacology (12.3)*]. Advise patients to avoid concomitant use with combined hormonal contraceptives (e.g., pill, patch, ring), implants, and progestin only pills. Counsel females using hormonal contraceptives to use

an alternative contraceptive method (e.g., non-hormonal intrauterine system) or additional non-hormonal contraceptive (e.g., condoms) during concomitant use and for 28 days after discontinuation of SKYCLARYS.

8.4 Pediatric Use

The safety and effectiveness of SKYCLARYS for the treatment of Friedreich's ataxia have been established in pediatric patients aged 16 years and older. Use of SKYCLARYS for this indication is supported by evidence from one adequate and well-controlled study (Study 1) in adults and in pediatric patients aged 16 years and older [see *Clinical Studies (14)*].

Safety and effectiveness of SKYCLARYS have not been established in pediatric patients less than 16 years of age.

8.5 Geriatric Use

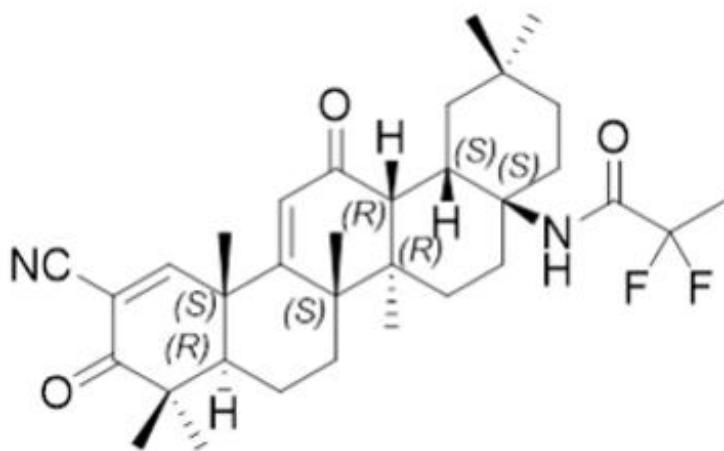
Clinical studies of SKYCLARYS in Friedreich's ataxia did not include patients aged 65 and over. No data are available to determine whether they respond differently than younger adult patients.

8.6 Hepatic Impairment

Omaveloxolone plasma exposure is increased in patients with moderate or severe hepatic impairment (Child-Pugh Class B and C) [see *Clinical Pharmacology (12.3)*]. Avoid treatment with SKYCLARYS in patients with severe hepatic impairment, including those who develop severe hepatic impairment. If hepatic function improves to moderate impairment, mild impairment, or normal function, initiation of SKYCLARYS treatment at the approved recommended dosage may be considered. For patients with moderate hepatic impairment, a reduced dosage is recommended with close monitoring for adverse reactions [see *Dosage and Administration (2.5)*]. For patients with mild hepatic impairment (Child-Pugh Class A), no dose adjustments are recommended .

11 DESCRIPTION

SKYCLARYS contains omaveloxolone in immediate release capsules for oral administration available in a 50 mg strength. The chemical name of omaveloxolone is N-(2-cyano-3,12-dioxo-28-noroleana-1,9(11)-dien-17-yl)-2,2-difluoro-propanamide. Omaveloxolone is a white to off-white amorphous solid. The molecular formula is $C_{33}H_{44}F_2N_2O_3$. Molecular weight is 554.72 g/mol. The chemical structure is:



Omaveloxolone has a pKa of 7.26 and is practically insoluble in water across the physiological pH range. Capsule contents include the following inactive ingredients: croscarmellose sodium, magnesium stearate, pregelatinized starch, and silicified microcrystalline cellulose. The hard capsule shells contain FD&C Blue #1, ferric oxide yellow, hypromellose, titanium dioxide, and white ink.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

The precise mechanism by which omaveloxolone exerts its therapeutic effect in patients with Friedreich's ataxia is unknown. Omaveloxolone have been shown to activate the Nuclear factor (erythroid-derived 2)-like 2 (Nrf2) pathway in vitro and in vivo in animals and humans. The Nrf2 pathway is involved in the cellular response to oxidative stress.

12.2 Pharmacodynamics

Potential to Prolong the QT Interval

The effect of omaveloxolone on the QTc interval has not been adequately characterized.

12.3 Pharmacokinetics

Absorption

The median (range) time to achieve peak plasma concentration was 7 to 14 (1 to 24) hours. The total plasma omaveloxolone exposure based on area under the concentration-time curve (AUC) increased in a dose-dependent and dose proportional manner over a dose range of 50 mg (0.33 times the recommended dosage) to 150 mg, but maximum omaveloxolone plasma concentration (C_{max}) increased in a less than dose proportional manner over the dose range in healthy fasted subjects.

Effect of Food

Omaveloxolone C_{max} and AUC_{0-inf} increased by approximately 350% and 15%, respectively, with a high-fat meal (800-1000 calories, approximately 150, 250, and 500 to 600 calories from protein, carbohydrate, and fat, respectively) compared to fasted conditions [see *Dosage and Administration* (2.2)].

Distribution

The mean apparent volume of distribution of omaveloxolone is 7361 L (105 L/kg for a 70 kg person). Protein binding of omaveloxolone is 97%.

Elimination

The mean (range) terminal half-life of omaveloxolone is 57 hours (32 to 90 hours). The mean apparent plasma clearance of omaveloxolone is 109 L/hr.

Metabolism

Omaveloxolone is primarily metabolized by CYP3A with minor metabolism by CYP2C8 and CYP2J2.

Excretion

Following administration of a single oral dose of radiolabeled omaveloxolone 150 mg to healthy subjects, approximately 92% of the dose was recovered in feces (approximately 91% within 96 hours after administration) and 0.1% in urine.

Specific Populations

There were no clinically significant differences in the pharmacokinetics of omaveloxolone based on age (16 to 71 years of age), sex, race, or body weight (41 to 128 kg). The effect of renal impairment on the pharmacokinetics of omaveloxolone is unknown.

Patients with Hepatic Impairment

There were no clinically significant differences in the pharmacokinetics of omaveloxolone in subjects with mild hepatic impairment (Child-Pugh Class A). In subjects with moderate and severe hepatic impairment (Child-Pugh Class B and C), omaveloxolone clearance was reduced, resulting in higher plasma exposure of omaveloxolone. The omaveloxolone AUC increased up to 1.65-fold and C_{\max} increased up to 1.83-fold in subjects with moderate hepatic impairment. The omaveloxolone AUC increased up to 2.17-fold in subjects with severe hepatic impairment; however, this change was variable [see *Use in Specific Populations* (8.6)].

Drug Interaction Studies

Clinical Studies

Strong CYP3A Inhibitors: Omaveloxolone C_{\max} increased 3-fold and AUC 4-fold following concomitant use with itraconazole (strong CYP3A inhibitor) [see *Drug Interactions* (7.1)].

Moderate CYP3A Inhibitors: Omaveloxolone C_{\max} and AUC increased approximately 1.25-fold following concomitant use with verapamil (moderate CYP3A4 and P-gp inhibitor) [see *Drug Interactions* (7.1)].

Strong and Moderate CYP3A Inducers: The effect of concomitant use with moderate and strong CYP3A4 inducers is unknown; however, a significant reduction in omaveloxolone exposure is likely following concomitant use based on its metabolic pathway.

Certain CYP450 Enzymes or Transporter Substrates: Omaveloxolone decreased the AUC of midazolam (CYP3A4 substrate) by approximately 45%, AUC of repaglinide (CYP2C8 substrate) by approximately 35%, and AUC of rosuvastatin (BCRP and OATP1B1 substrate) by approximately 30% [see *Drug Interactions* (7.2)]. There were no clinically

significant differences in the pharmacokinetics of digoxin (P-gp substrate) or metformin [(organic cation transporter (OCT)1 substrate] when co-administered with omaveloxolone.

Other Drugs: No clinically significant differences in the pharmacokinetics of omaveloxolone are expected following concomitant use with weak CYP3A4 inhibitors or strong CYP2C8 inhibitors.

In Vitro Studies

CYP Enzymes: Omaveloxolone is not an inhibitor of CYP1A2, CYP2B6, CYP2C9, CYP2C19, and CYP2D6. Omaveloxolone is not an inducer of CYP1A2 and CYP2B6.

Drug Transporters: Omaveloxolone is not an inhibitor of BCRP, BSEP, OAT3, OATP1B1, OATP1B3, OCT2, MATE1, and MATE2-K. Omaveloxolone inhibited the renal transporter OAT1.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis

Carcinogenicity studies have not been conducted with omaveloxolone.

Mutagenesis

Omaveloxolone was negative in a bacterial reverse mutation (Ames) assay, and positive in a chromosomal aberration assay in human peripheral blood lymphocytes but negative in in vitro (rat micronucleus and comet) assays.

Impairment of Fertility

Oral administration of omaveloxolone (0, 1, 3, and 10 mg/kg/day) to male and female rats prior to and during mating and continuing in females to gestation day 7 produced an increase in pre- and post-implantation loss and resorptions, resulting in a decrease in viable embryos at the highest dose tested. The no-effect dose (3 mg/kg/day) for adverse effects on fertility and reproductive function was associated with plasma exposures (AUC) approximately 2 times that in humans at the recommended human dose of 150 mg/day.

14 CLINICAL STUDIES

The efficacy of SKYCLARYS was evaluated in a 48-week, randomized, double-blind, placebo-controlled study in patients 16 to 40 years of age with Friedreich's Ataxia (Study 1; NCT02255435).

A total of 103 patients were randomized (1:1) to receive SKYCLARYS 150 mg once daily (n=51) or placebo (n=52).

Enrolled patients had to have a stable modified Friedreich's Ataxia Rating Scale (mFARS) score between 20 and 80, be able to complete maximal exercise testing, and have a left ventricular ejection fraction of at least 40%. In Study 1, 53% of enrolled patients were male, 97% were White, and the mean age was 24 years at study entry. Patients with or

without pes cavus were included in Study 1. Pes cavus was defined as having a loss of lateral support and was determined if light from a flashlight could be seen under the patient's arch when barefoot and weight bearing.

The prespecified primary analysis was the change from baseline in the mFARS score compared to placebo at Week 48 in the Full Analysis Population of patients without pes cavus (n=82). The mFARS is a clinical assessment tool to assess patient function, which consists of 4 domains to evaluate bulbar function, upper limb coordination, lower limb coordination, and upright stability. The mFARS has a maximum score of 99, with a lower score on the mFARS signifying lesser physical impairment.

Treatment with SKYCLARYS resulted in statistically significant lower mFARS scores (less impairment) relative to placebo (see Table 4) at Week 48.

Table 4: Primary Analysis in Full Analysis Population: mFARS Least Squares (LS) Mean Change from Baseline at Week 48

	Mean (SD) Baseline mFARS Total Score	LS Mean Change from Baseline at Week 48	Treatment difference SKYCLARYS- placebo (95% CI)	p-value
SKYCLARYS (n = 40)	40.95 (10.39)	-1.56	-2.41 (-4.32, -0.51)	0.0138
Placebo (n = 42)	38.78 (11.03)	0.85		

SD = Standard Deviation; LS = Least Squares; CI = Confidence Interval

The All Randomized Population (N=103), which included all patients regardless of pes cavus status, demonstrated similar results to the Full Analysis Population of lower mFARS scores in patients treated with SKYCLARYS compared to placebo, with a nominally significant least squares mean difference between treatment groups of -1.94 (95% CI: -3.71, -0.16, p=0.0331).

In a post hoc, propensity-matched analysis, lower mFARS scores were observed in patients treated with SKYCLARYS after 3 years relative to a matched set of untreated patients from a natural history study. These exploratory analyses should be interpreted cautiously given the limitations of data collected outside of a controlled study, which may be subject to confounding.

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

SKYCLARYS (omaveloxolone) capsules, 50 mg, are supplied as opaque, hard capsules having a light green body and blue cap imprinted with "RTA 408" in white ink on the body and "50" in white ink on the cap. SKYCLARYS is supplied in high density polyethylene bottles that contain 90 capsules, with a foil induction seal and child-resistant closure (NDC 73179-250-90).

16.2 Storage and Handling

Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C and 30°C (59°F to 86°F) [see USP Controlled Room Temperature].

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Patient Information).

Elevation of Aminotransferases

Inform patients that elevation in aminotransferases have occurred in patients treated with SKYCLARYS. Liver function tests will be performed prior to initiating SKYCLARYS, every month for the first 3 months of treatment, and periodically thereafter as needed [see *Warnings and Precautions (5.1)*].

Fluid Overload

Inform patients that elevations in BNP, a marker of cardiac function, have occurred in patients treated with SKYCLARYS. BNP will be performed prior to initiating SKYCLARYS and if signs and symptoms of fluid overload occur, such as sudden weight gain, peripheral edema, palpitations, and shortness of breath. Advise patients to contact their healthcare provider if signs and symptoms of fluid overload develop [see *Warnings and Precautions (5.2)*].

Lipid Abnormalities

Inform patients that treatment with SKYCLARYS has been associated with increases in LDL cholesterol and decreases in HDL cholesterol. Cholesterol will be assessed prior to starting SKYCLARYS and monitored periodically during treatment [see *Warnings and Precautions (5.3)*].

Drug Interactions

Advise patients to discuss all medications they are taking, including other prescription medications, non-prescription medications, or herbal products (e.g., St. John's Wort) with their healthcare provider [see *Drug Interactions (7)*].

Pregnancy

Advise women to notify their healthcare provider if they become pregnant or intend to become pregnant during SKYCLARYS therapy [see *Use in Specific Populations (8.1)*].

Females of Reproductive Potential

Counsel females using hormonal contraceptives to use an alternative contraceptive method (e.g., non-hormonal intrauterine system) or additional non-hormonal contraceptive (e.g., condoms) during concomitant use and for 28 days after discontinuation of SKYCLARYS [see *Drug Interactions (7.2)* and *Use in Specific Populations (8.3)*].

Administration

Advise patients to take SKYCLARYS on an empty stomach at least 1 hour before eating [see *Dosage and Administration (2.2)*].

Swallow SKYCLARYS capsules whole. Do not open, crush, or chew.

Advise patients that if a dose of SKYCLARYS is missed to not to double their dose or take more than the prescribed dose [see *Dosage and Administration* (2.3)].

Advise patients to avoid grapefruit juice and grapefruit while they are taking SKYCLARYS [see *Drug Interactions* (7.1)].

Manufactured for Reata Pharmaceuticals, Inc., Plano, TX 75024 USA

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PATIENT INFORMATION
SKYCLARYS™ (skye klar' is)
(omaveloxolone)
capsules, for oral use

What is SKYCLARYS?

SKYCLARYS is used for the treatment of Friedreich's ataxia in adults and children 16 years of age and older.

It is not known if SKYCLARYS is safe and effective for use in children younger than 16 years of age.

Before taking SKYCLARYS, tell your healthcare provider about all of your medical conditions, including if you:

- have liver problems.
- have a history of heart problems, including heart failure.
- have a high level of fat in your blood (high blood cholesterol).
- are pregnant or plan to become pregnant.
- are breastfeeding or plan to breastfeed. It is not known if SKYCLARYS passes into your breast milk. Talk to your healthcare provider about the best way to feed your baby if you take SKYCLARYS.
- It is not known if SKYCLARYS will harm your unborn baby.
- Women who use hormonal birth control should use another form of birth control such as a non-hormonal intrauterine system or an extra non-hormonal birth control such as condoms while using SKYCLARYS and for 28 days after stopping SKYCLARYS.

Tell you healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements such as St. John's Wort.

Taking SKYCLARYS with other medicines can cause serious side effects.

SKYCLARYS may affect the way other medicines work, and other medicines may affect how SKYCLARYS works.

Know the medicines you take. Keep a list of them to show to your healthcare provider and pharmacist when you get a new medicine.

How should I take SKYCLARYS?

- Take SKYCLARYS exactly as your healthcare provider tells you to take it.

- Take SKYCLARYS capsules on an empty stomach at least 1 hour before eating.
- Swallow SKYCLARYS capsules whole. Do not open, crush, or chew.
- If you miss a dose, then you should skip the missed dose and take the next dose at the regular time the next day. Do not double your next dose or take more than the prescribed dose.

What should I avoid while taking SKYCLARYS?

Do not drink grapefruit juice or eat grapefruit. These may change the amount of SKYCLARYS in your blood.

What are the possible side effects of SKYCLARYS?

SKYCLARYS may cause serious side effects, including:

- **increase in blood liver enzymes.** Some people taking SKYCLARYS have had an increase in the level of liver enzymes in their blood. Your healthcare provider will do liver function tests

- before you start taking SKYCLARYS
- every month for the first 3 months after starting your treatment with SKYCLARYS

SKYCLARYS

- during certain times as needed while taking SKYCLARYS

If your liver enzymes increase, your healthcare provider may change your dose during treatment, stop treatment for some time, or completely stop treatment with SKYCLARYS.

- **increase in a blood protein called B-Type Natriuretic Peptide (BNP).** BNP tells how well your heart is working. Your healthcare provider will check your BNP levels before your treatment with SKYCLARYS. Tell your healthcare provider if you have signs and symptoms of your heart not working well such as too much fluid in your body (fluid overload). Signs and symptoms may include:

- sudden weight gain (3 pounds or more of weight gain in 1 day, or 5 pounds or more of weight gain in 1 week)
- swelling in your arms, hands, legs, or feet (peripheral edema)
- fast heartbeat (palpitations)
- shortness of breath

If you have symptoms of fluid overload that is considered a side effect of SKYCLARYS, your healthcare provider may stop treatment with SKYCLARYS.

- **changes in cholesterol levels.** Increases in low density lipoprotein cholesterol (LDL-C) or bad cholesterol and decreases in high density lipoprotein cholesterol (HDL-C) or good cholesterol have happened during treatment with SKYCLARYS.

Your healthcare provider will check your cholesterol levels before and during your treatment with SKYCLARYS.

The most common side effects of SKYCLARYS include:

- increased liver enzymes (ALT/AST)
- headache
- nausea
- stomach pain
- tiredness

- diarrhea
- muscle pain

These are not all the possible side effects of SKYCLARYS.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store SKYCLARYS?

- Store SKYCLARYS at room temperature between 68°F to 77°F (20°C to 25°C)

Keep SKYCLARYS and all medicines out of the reach of children.

General information about the safe and effective use of SKYCLARYS.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use SKYCLARYS for a condition for which it was not prescribed. Do not give SKYCLARYS to other people, even if they have the same symptoms you have. It may harm them. You can ask your healthcare provider or pharmacist for information about SKYCLARYS that is written for health professionals.

What are the ingredients in SKYCLARYS?

Active ingredient: omaveloxolone

Inactive ingredients: croscarmellose sodium, magnesium stearate, pregelatinized starch, silicified microcrystalline cellulose. The hard capsule shells contain FD&C Blue #1, ferric oxide yellow, hypromellose, titanium dioxide, and white edible ink.

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For more information about SKYCLARYS™, go to www.SKYCLARYS.com or call Reata Pharmaceuticals, Inc. at 1-800-314-3934.

This Patient Information has been approved by the U.S. Food and Drug Administration
Approved: 2/2023

PRINCIPAL DISPLAY PANEL

73179-250-90

SKYCLARYS

(omaveloxolone)

50 mg capsules

Rx only

90 capsules

SKU FPO
1 234567 890128 >

Recommended Dosage: Take 150 mg (3 capsules) orally once daily. Each capsule contains 50 mg of omeveloxolone.

Swallow SKYCLARYS capsules whole. Do not open, crush, or chew. Keep out of reach of children.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C and 30°C (59°F and 86°F) [see USP Controlled Room Temperature]. (800) 314-3394

73179-250-90

SKYCLARYS™
(omeveloxolone) capsules

50 mg

TA

50

Rx Only
90 capsules

REATA
PHARMACEUTICALS

Manufactured for Reata Pharmaceuticals, Inc., Plano, TX 75024 USA
by Parthenon Pharmaceuticals, Inc., Cincinnati, OH 45237 USA

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SN 123456789012
EXP YYYY/MMMM
Lot ABC123

FPO

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SKYCLARYS

omeveloxolone capsule

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:73179-250
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
OMAVELOXOLONE (UNII: G69Z98951Q) (OMAVELOXOLONE - UNII:G69Z98951Q)		OMAVELOXOLONE	50 mg
Inactive Ingredients			
Ingredient Name			Strength
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)			
Product Characteristics			
Color	green (light green body and blue cap)	Score	no score
Shape	CAPSULE	Size	22mm
Flavor		Imprint Code	RTA;408;50
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73179-250-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/28/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA216718	02/28/2023	

Labeler - Reata Pharmaceuticals, Inc. (128566457)

Revised: 5/2023

Reata Pharmaceuticals, Inc.